PATENT

Attorney Docket No.: JHU1290-7

In re Application of: James E. Hildreth

. Application No.: 09/761,209 Filed: January 16, 2001

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Amendments to the Claims

Please amend claims 8, 11, 12, 24-27, 29, and 31, as indicated in the Listing of Claims.

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

Claims 1-7 (Canceled)

8. (Currently Amended): A method of ameliorating an autoimmune disease or graft

rejection in an animal, the method comprising administering to the animal a therapeutically

effective amount of an a monoclonal antibody, capable of suppressing intercellular leukocyte-

leukocyte adhesion, wherein the antibody binds to an epitope on the leukocyte adhesion receptor

 β -chain, thereby ameliorating the autoimmune disease or graft rejection in the animal.

9. (Original): The method of claim 8, wherein the receptor is selected from the group

consisting of LFA-1, Mac-1, and Leu M5.

10. (Canceled)

11. (Currently Amended): The method of claim 8, wherein the monoclonal antibody has

the specificity of the monoclonal antibody produced by ATCC HB [[X]] 10160.

12. (Currently Amended): The method of claim 8, wherein the antibody is produced by

hybridoma cell line ATCC HB [[X]] 10160.

13. (Original): The method of claim 8, wherein the administration is parenteral.

14. (Original): The method of claim 13, wherein the parenteral administration is by

subcutaneous, intramuscular, intraperitoneal, intracavity, transdermal, or intravenous injection.

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- 15. (Original): The method of claim 8, wherein said administration is at a dosage of about 0.01 mg/kg/dose to about 2000 mg/kg/dose.
- 16. (Original): The method of claim 8, wherein the monoclonal antibody is therapeutically labeled.
- 17. (Original): The method of claim 16, wherein the therapeutic label is selected from the group consisting of a radioisotope, a drug, a lectin, and a toxin.

Claims 18-23 (Canceled)

- 24. (Currently Amended): A method of ameliorating acquired immunodeficiency syndrome (AIDS), an antoimmune disease, or graft rejection in an animal, suppressing HIV-induced cell fusion, comprising: administering to the animal a therapeutically contacting a leukocyte infected with HIV with an effective amount of an antibody, capable of suppressing intercellular leukocyte-leukocyte adhesion, wherein the antibody binds to an epitope on the leukocyte adhesion receptor β -chain, thereby ameliorating acquired immunodeficiency syndrome (AIDS), an autoimmune disease, or graft rejection in the animal suppressing HIV-induced cell fusion.
- 25. (Currently Amended): The method of claim 24, wherein the receptor is selected from the group consisting of LFA-1, Mac-1, and Leu M5.
- 26. (Currently Amended): The method of claim 24, wherein the monoclonal antibody has the specificity of the monoclonal antibody produced by ATCC HB [[X]] 10160.

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27. (Currently Amended): The method of claim 24, wherein the antibody said administration is is administered at a dosage of about 0.01 mg/kg/dose to about 2000 mg/kg/dose.

- 28. (Previously Presented): The method of claim 24, wherein the monoclonal antibody is therapeutically labeled.
- 29. (Currently Amended): A method of ameliorating graft rejection in an animal, the method comprising administering to the animal a therapeutically effective amount of an antibody, capable of suppressing intercellular leukocyte-leukocyte adhesion, wherein the antibody binds to an epitope on the leukocyte adhesion receptor β -chain, thereby ameliorating the graft rejection in the patient.
- 30. (Previously Presented): The method of claim 29, wherein the receptor is selected from the group consisting of LFA-1, Mac-1, and Leu M5.
- 31. (Currently Amended): The method of claim 29, wherein the antibody has the specificity of the monoclonal antibody produced by ATCC HB [[X]] 10160.
- 32. (Previously Presented): The method of claim 29, wherein said administration is at a dosage of about 0.01 mg/kg/dose to about 2000 mg/kg/dose.
- 33. (Previously Presented): The method of claim 29, wherein the monoclonal antibody is therapeutically labeled.
 - 34. (Previously Presented): The method of claim 29, wherein the animal is a human.